Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU)

COMMISSION IMPLEMENTING DECISION

of 17 February 2012

amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres

(notified under document C(2012) 860)

(Text with EEA relevance)

(2012/112/EU)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC⁽¹⁾, and in particular the first paragraph of Article 22 thereof,

Whereas:

- (1) Directive 92/65/EEC lays down the animal health requirements governing trade in the Union in animals, semen, ova and embryos not subject to the animal health requirements laid down in certain specific Union acts. In addition, Part 1 of Annex E to that Directive sets out the specimen health certificate for trade in animals from holdings (ungulates, birds, lagomorphs, dogs, cats and ferrets), while Part 3 of that Annex sets out the specimen health certificate for trade in animals, semen, embryos and ova from approved bodies, institutes or centres.
- (2) Article 6(3) of Directive 92/65/EEC lays down the animal health requirements governing trade in suidae other than those covered by Council Directive 64/432/EEC of 26 June 1964 on animal health problems affecting intra-Community trade in bovine animals and swine⁽²⁾. It provides, inter alia, that where suidae do not come from a brucellosis-free herd in accordance with Directive 64/432/EEC, they must, in the 30 days prior to their dispatch, have undergone with negative results a test designed to show the absence of antibodies to brucellosis. In the interests of consistency of Union legislation, the specimen health certificate set out in Part 1 of Annex E to

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

Directive 92/65/EEC should therefore be amended to include a specific reference to that requirement.

- (3) Commission Decision 2007/598/EC of 28 August 2007 concerning measures to prevent the spread of highly pathogenic avian influenza to other captive birds kept in zoos and approved bodies, institutes or centres in the Member States⁽³⁾ approves preventive vaccination plans against that disease in certain Member States.
- (4) Point 4(b) of Annex II to Decision 2007/598/EC provides that birds vaccinated against avian influenza kept in zoos that are not approved in accordance with Directive 92/65/ EEC may be moved to other Member States, after authorisation by the Member State of destination, provided that they meet the requirements set out in that Decision and they are accompanied by a health certificate, as laid down in Part 1 of Annex E to that Directive, specifying that they are conform to Decision 2007/598/EC and are vaccinated against avian influenza on a specified date.
- (5) However, birds as referred to in Article 7 of Directive 92/65/EEC are not required to be accompanied by a health certificate, as set out in Part 1 of Annex E thereto when traded within the Union, but must be accompanied by a self-certification by the operator in accordance with Article 4 of that Directive, or in the case of psittacidae by a commercial document signed by the official veterinarian or by the veterinarian responsible for the holding.
- (6) It should be therefore clarified that the health certificate set out in Part 1 of Annex E to Directive 92/65/EEC is only required to accompany birds that are vaccinated against avian influenza and come from a holding on which vaccination against avian influenza was carried out during the past 12 months. Therefore, the specimen health certificate set out in Part 1 of that Annex should be amended to include a reference to such vaccination.
- (7) Article 10 of Directive 92/65/EEC lays down the animal health requirements governing trade in dogs, cats and ferrets. It provides, inter alia, that they must satisfy the relevant requirements laid down in Regulation (EC) No 998/2003 of the European Parliament and of the Council of 26 May 2003 on the animal health requirements applicable to the non-commercial movement of pet animals and amending Council Directive 92/65/EEC⁽⁴⁾.
- (8) Article 6 of Regulation (EC) No 998/2003 provides that until 31 December 2011, dogs and cats entering Ireland, Malta, Sweden and the United Kingdom from other Member States are to be vaccinated and subject to a pre-entry rabies blood testing in accordance with national rules.
- (9) In addition, Article 16 of that Regulation provides that until 31 December 2011, Finland, Ireland, Malta, Sweden and the United Kingdom, as regards echinococcosis, and Ireland, Malta and the United Kingdom as regards ticks, may make the entry of pet animals into their territory subject to compliance with certain additional national requirements.
- (10) Commission Delegated Regulation (EU) No 1152/2011 of 14 July 2011 supplementing Regulation (EC) No 998/2003 of the European Parliament and of the Council as regards preventive health measures for the control of *Echinococcus multilocularis* infection in

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

dogs⁽⁵⁾ was adopted in order to ensure the continuous health protection of Ireland, Malta, Finland and the United Kingdom from *Echinococcus multilocularis*. It is to apply from 1 January 2012.

- (11) The reference to Articles 6 and 16 of Regulation (EC) No 998/2003 included in the specimen health certificate set out in Part 1 of Annex E to Directive 92/65/EEC should therefore be deleted and replaced, as regards dogs, by a reference to Delegated Regulation (EU) No 1152/2011.
- (12) Part 1 of Annex E to Directive 92/65/EEC should therefore be amended accordingly.
- (13) Article 13 of Directive 92/65/EEC lays down the animal health requirements governing trade in animals of species susceptible to the diseases listed in Annexes A and B thereto and in semen, ova and embryos of such animals consigned to and from bodies, institutes or centres approved in accordance with Annex C thereto.
- (14) Semen, ova and embryos of certain animal species can be frozen and stored for a long time and therefore donor animal might no longer be available on the day the health certificate is issued. It is therefore necessary to amend the specimen health certificate set out in Part 3 of Annex E to Directive 92/65/EEC to state that the donor animal was found to be healthy and free from clinical disease either on day of collection or the date of issuing of the health certificate.
- (15) Point 4(a) of Annex II to Decision 2007/598/EC provides that birds vaccinated against avian influenza kept in approved bodies, institutes or centres including zoos may only be moved to approved bodies, institutes or centres including zoos in other Member States provided that they meet the requirements set out in that Decision and they are accompanied by a health certificate as laid down in Part 3 of Annex E to Directive 92/65/ EEC stating that the birds have been vaccinated against avian influenza in conformity to Commission Decision 2006/474/EC⁽⁶⁾. As that Decision has since been repealed and replaced by Decision 2007/598/EC, that reference should be replaced by a reference to Decision 2007/598/EC.
- (16) Part 3 of Annex E to Directive 92/65/EEC should therefore be amended accordingly.
- (17) Directive 92/65/EEC should therefore be amended accordingly.
- (18) To avoid any disruption of trade, the use of health certificates issued in accordance with Part 1 and Part 3 of Annex E to Directive 92/65/EEC, before the amendments introduced by this Decision, should be authorised during a transitional period subject to certain conditions.
- (19) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

Annex E to Directive 92/65/EEC is amended in accordance with the Annex to this Decision.

Status: Point in time view as at 31/01/2020. Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

Article 2

For a transitional period until 30 June 2012, Member States may authorise trade in animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres accompanied by a health certificate issued not later than 29 February 2012 in accordance with the models set out in Part 1 and Part 3 of Annex E to Directive 92/65/EEC in its version prior to the amendments introduced by this Decision.

Article 3

This Decision shall apply from 1 March 2012.

Article 4

This Decision is addressed to the Member States.

Done at Brussels, 17 February 2012.

For the Commission John DALLI Member of the Commission

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

ANNEX

Annex E to Directive 92/65/EEC is amended as follows:

(1) Part 1 is replaced by the following:

Part 1 —

Health certificate for trade in animals from holdings (ungulates, birds vaccinated against avian influenza, lagomorphs, dogs, cats and ferrets) 92/65 EI

ROPE	ΞA	N UNION	Intra trade certifica		
1.1		Consignor Name	I.2. Certificate reference No I.2.a. Local reference No		
		Address Postal code	I.3. Central competent authority		
			I.4. Local competent authority		
1.5		Consignee Name	I.6. No(s) of related original certificates documents		
		Address Postal code	1.7.		
I.5 I.8		Country of origin ISO code I.9. Region of Code origin	I.10. Country of ISO code I.11. Region of Code destination		
1.1	2.	Place of origin Holding	I.13. Place of destination Holding Establishment Approved body		
		Name Approval number Address	Name Approval number Address		
		Postal code	Postal code		
1.1	4.	Place of loading Postal code	I.15. Date and time of departure		
1.1	6.	Means of transport	I.17. Transporter		
		Aeroplane Bip Railway wagon Road vehicle Other I	Name Approval number Address		
		Identification	Postal code		
1.1	I.18. Description of commodity I.19. Commodity code (CN code)				
			I.20. Quantity		
1.2	1.		I.22. Number of packages		
1.2	3.	Seal/Container No	1.24.		
1.2	5.	Commodities certified for:			
		Breeding Production Artificial reproduction	Slaughter Pets Approved body		
1.2	6.	Transit through third country	I.27. Transit through Member States		
		Third country ISO code Exit point Code	Member State ISO code Member State ISO code		
	_	Entry point BIP No	Member State ISO code		
1.2	8.	Export Third country ISO code Exit point Code	I.29. Estimated journey time		
1.3	0.	Route plan Yes No	1		
12	1	Identification of the commodities			
1.3			ation number Sex Age Quantity		

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

EUROPEAN UNION				92/65 El Animals from holdings (ungulates, birds (²), lagomorphs, dogs, cats and ferrets)			
П.		Health	information	II.a. Certificate reference number	II.b.		
			indersigned official veterinarian (1)/veterinarian responsible γ (1) certify that:	onsible for the establishment of origin	n and approved by the competer		
(¹) eiti	her	(II.1.	at the time of inspection the above animals wer provisions of Council Regulation (EC) No 1/2005.]		ed journey in accordance with th		
(¹) or		[.1.	at the time of inspection the dogs (1)/cats (1)/ferrets (1) to be moved for non-commercial purposes in accordance with Commission Regulation (EU) No 388/2010 were fit to travel.]				
(¹) eiti	her	[II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the ruminant(s) (1)/suidae (1) other than that/those covered by Council Directive 64/432/EEC (1) or Council Directive 91/68/EEC (1):				
			(a) belong(s) to the species				
			(b) at the time of examination, do(does) not show	any clinical sign of any disease to wi	hich it/they is/are susceptible;		
			(c) come(s) from an officially tuberculosis-free (¹) subject to swine fever restrictions or from a he laid down in Article 6(2)(b) (¹)(the test laid down	olding where it/they was/were subjected	d with negative results to the tes		
(¹) (²)	or	[II.2.	the conditions of Article 4 of Council Directive 92/ Directive 2009/158/EC:	65/EEC are fulfilled and the birds othe	er than those referred to in Coun		
			 (a) conform to Decision 2007/598/EC and were vavaccine				
			(b) satisfy the requirements of Article 7 of Council	Directive 92/65/EEC;			
			(c) at the time of examination do not show any cl	inical sign of any disease to which the	ey are susceptible.]		
(1) or		[11.2.	the conditions of Article 4 of Council Directive 92/	65/EEC are fulfilled and the lagomorp	hs:		
			(a) satisfy the requirements of Article 9 of Counci	Directive 92/65/EEC;			
			(b) at the time of examination do not show any cl	inical signs of disease to which they a	are susceptible.]		
(¹) or		[II.2.	the conditions of Article 4 of Council Directive 92/66 hours before dispatch, by a veterinarian authorisec be in good health, and satisfy, in accordance with / Article 5 of Regulation (EC) No 998/2003 of the E	by the competent authority, and that Article 10(2) of Council Directive 92/65/	examination showed the animals EEC, the requirements laid down		
and	(1)	either	[have not been treated against Echinococcus mult	ilocularis.]			
	(¹)	or	[have been treated against <i>Echinococcus multi</i> No 1152/2011.]]	<i>locularis</i> in accordance with Comm	ission Delegated Regulation (E		
(¹) or		[II.2.	the conditions of Article 4 of Council Directive 92/6 ation, within 24 hours before dispatch, by a veterin the animals to be in good health, and satisfy, requirements laid down in Article 5 of Regulation	arian authorised by the competent auth in accordance with Article 10(2) of	nority, and that examination show Council Directive 92/65/EEC. t		
(¹) or		[11.2.	the consignment of more than five dogs to be mo lation (EU) No 388/2010 underwent a clinical exar the competent authority, and that examination s accordance with Article 10(2) of Council Directive No 998/2003 of the European Parliament and of t	nination, within 24 hours before dispat howed the animals to be in good h 9 92/65/EEC, the requirements laid do	tch, by a veterinarian authorised nealth, and the animals satisfy,		
and	(1)	either	[their scheduled destination indicated in Box I.10, c against <i>Echinococcus multilocularis</i> in accordance				
	(¹)	or	[they have been treated against <i>Echinococcus mu</i> lation (EU) No 1152/2011.]]	Iltilocularis in accordance with Article	7 of Commission Delegated Reg		

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

EUROPEAN UNION			92/65 El Animals from holdings (ungulates, birds (²), lagomorph dogs, cats and ferrets)		
II.	Health information		II.a. Certificate reference number	II.b.	
(¹) or	Regulation (EU) No 3 the competent author with Article 10(2)	888/2010 underwent a clinical examity, and that examination showed	be moved for non-commercial purpos nination, within 24 hours before dispa the animals to be in good health and D, the requirements laid down in puncil;]	tch, by a veterinarian authorised I I the animals satisfy, in accordance	
II.3.	The additional guarantees regarding diseases listed in Annex B (3) to Council Directive 92/65/EEC are as follows: (1)				
	Disease	Decision			
	Disease	Decision			
	Disease	Decision			
II.4.	This certificate is valid until	(4)			
Notes					
Part I:					
— Box r	references I.1 to I.4, I.8, I.20, I.25	and I.31: Required for non-comme	ercial movement of more than five do	gs, cats and ferrets.	
— Box r	reference I.6: No(s) of accompar	nying documents: CITES, if applica	able.		
— Box r	reference I.19: Use the appropriat	e HS code: 01.06.19, 01.06.31, 0	1.06.32, 01.06.39.		
— Box r	reference I.25: Indicate "Pets" on	ly when more than five dogs, cal	ts or ferrets are to be certified for s	strictly non-commercial movemen	
— Box r	reference I.31: Identification syste cation may be use		used wherever possible but in the c	ase of small animals, batch ident	
Part II:					
(1) Delet	te as necessary.				
	fication requirements only apply to mission Decision 2007/598/EC.	birds that have been vaccinated	against avian influenza under a preve	entive vaccination plan approved	
		efiting from additional guarantees u	under Union legislation.		
(⁴) The paccol	period of validity of this certificate rdance with Commission Regulatio	is 10 days from the date of issue,	except for dogs, cats and ferrets more the certificate is valid for a period of ferrets		
— The d	colour of the stamp and signature	must be different from that of the	other particulars in the certificate.		
	veterinarian or official inspector				
Official v					
	me (in capital letters):		Qualification and title:		
Na	me (in capital letters): cal veterinary unit:		Qualification and title: LVU No:		
Na	cal veterinary unit:				
Na Loc Dal	cal veterinary unit:		LVU No:		

(2) Part 3 is replaced by the following:

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

Part 3 —

Health certificate for trade in animals, semen, ova and embryos from approved bodies, institutes or centres 92/65 EIII EUROPEAN UNION

EUR	OPEAN UNION Intra trade certificat					
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No			
Ited		Address	I.3. Central competent authority			
		Postal code	I.4. Local competent authority			
t presen	1.5.	Consignee Name	I.6. No(s) of related original certificates documents			
gnment		Address Postal code	l.7.			
of consignment presented	1.8.	Country of origin ISO code I.9. Region of Code origin	I.10. Country of ISO code I.11. Region of Code destination			
Part I: Details	l.12.	Place of origin Approved body	I.13. Place of destination Approved body			
Part I:		Name Approval number Address	Name Approval number Address			
		Postal code	Postal code			
	I.14.	Place of loading Postal code	I.15. Date and time of departure			
	I.16.	Means of transport	I.17. Transporter			
		Aeroplane Ship Railway wagon Railway wagon I Road vehicle Other I Identification	Name Approval number Address			
			Postal code			
	l.18.	Description of commodity	I.19. Commodity code (CN code)			
			I.20. Quantity			
	I.21.		I.22. Number of packages			
	1.23.	Seal/Container No	1.24.			
	1.25.	Commodities certified for:				
		Approved body				
	1.26.	Transit through third country	I.27. Transit through Member States			
		Third country ISO code	Member State ISO code			
		Exit point Code Entry point BIP No	Member State ISO code Member State ISO code			
	1.28	Export	I.29. Estimated journey time			
		Third country ISO code				
		Exit point Code				
	1.30.	Route plan				
		Yes No				
	1.31.	Identification of the commodities				
	Species Identification system Identification number Sex Age Quantity (scientific name)					

Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

П.	OPEAN UNION		92/65 EIII Animals from approved bodies, institutes or centres			
L	Health inform	ation	II.a. Certificate reference number	II.b.		
	I, the undersination authority (¹) ce	gned official veterinarian (¹)/veterinarian respons rtify that:	ible for the establishment of origin a	and approved by the compete		
II.1.	 The body, institute or centre of origin is approved in accordance with Annex C to Council Directive 92/65/EEC for the purpose of trade the animals, semen, ova or embryos described in Box I.18. 					
 II.2. The animals, semen, ova or embryos described in Box I.18. II.2. The animals (1)/donor animals (1) described in this certificate have been examined today (1)/on the day of collection (1) and found healthy and free of clinical signs of infectious diseases including those listed in Annex A to Directive 92/65/EEC and are not subject of official restrictions and remained in this body, institute or centre either since birth or for the following time						
11.3.		nspection, the above animals were fit to be transp C) No 1/2005 and IATA requirements and/or CITE				
II.4. The additional guarantees regarding diseases listed in Annex B (²) to Council Directive 92/65/EEC are as follows: (¹)						
	Disease	Decision	()			
	Disease	Decision				
{	Disease	Decision				
	Diotaco					
[II.5.		ing to Decision 2007/598/EC were vaccinated aga n an approved body, institute or centre of origin on)				
Note	95					
Part	art I:					
- Box reference I.6: No(s) of accompanying documents: CITES, if applicable.						
- e	Box reference I.19:	Use the appropriate HS code: 01.06.11, 01.06.1	9, 01.06.31, 01.06.32, 01.06.39, 05.11	.99.85.		
- B	Box reference I.31:	Identification system: individual identification mu identification may be used.	ist be used wherever possible but in	the case of small animals, bate		
In the case of semen, ova and embryos it indicated in the following format: official id						
		indicated in the following format: official identification	ation of the animal/dd/mm/yyyy.	he date of collection and shall b		
		indicated in the following format: official identification Age and sex: to be completed only in the case	ation of the animal/dd/mm/yyyy. of live animals, if appropriate.			
		indicated in the following format: official identification	ation of the animal/dd/mm/yyyy. of live animals, if appropriate.			
Part	11:	indicated in the following format: official identifica Age and sex: to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryo	ation of the animal/dd/mm/yyyy. of live animals, if appropriate.			
		indicated in the following format: official identific: <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated.	ation of the animal/dd/mm/yyyy. of live animals, if appropriate.			
(¹) [Delete as necessar	indicated in the following format: official identifica <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated. y.	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or			
(¹) C (²) A	Delete as necessar As requested by a	indicated in the following format: official identific: <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated.	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or ees under Union legislation.	other packaging express as uni		
(¹) C (²) A — T	Delete as necessar As requested by a	indicated in the following format: official identifica <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated. y. Member State benefiting from additional guarante tamp and signature must be different from that of	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or ees under Union legislation.	other packaging express as uni		
(¹) C (²) A — T	Delete as necessar As requested by a The colour of the st	indicated in the following format: official identifica <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated. y. Member State benefiting from additional guarante tamp and signature must be different from that of official inspector	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or ees under Union legislation.	other packaging express as un		
(¹) C (²) A — T	Delete as necessar As requested by a The colour of the s cial veterinarian or	indicated in the following format: official identifica <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated. y. Member State benefiting from additional guarante tamp and signature must be different from that of official inspector letters):	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or ees under Union legislation. f the other particulars in the certificate.	other packaging express as uni		
(¹) C (²) A — T	Delete as necessar As requested by a The colour of the si ial veterinarian or Name (in capital	indicated in the following format: official identifica <i>Age</i> and <i>sex</i> : to be completed only in the case <i>Quantity</i> : in the case of semen, ova and embryor should be indicated. y. Member State benefiting from additional guarante tamp and signature must be different from that of official inspector letters):	ation of the animal/dd/mm/yyyy. of live animals, if appropriate. os the number of straws, ampoules or ees under Union legislation. f the other particulars in the certificate. Qualification and title:	other packaging express as uni		
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Changes to legislation: Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

- (1) OJ L 268, 14.9.1992, p. 54.
- (2) OJ 121, 29.7.1964, p. 1977/64.
- (**3**) OJ L 230, 1.9.2007, p. 20.
- (4) OJ L 146, 13.6.2003, p. 1.
- (5) OJ L 296, 15.11.2011, p. 6.
- (6) OJ L 187, 8.7.2006, p. 37.

Status:

Point in time view as at 31/01/2020.

Changes to legislation:

Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 12 December 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations.